

Bend, OR, KBDN, VOR RWY 16, Amdt 11  
Florence, SC, KFLO, ILS OR LOC RWY 9,  
Amdt 13  
Florence, SC, KFLO, RNAV (GPS) RWY 1,  
Amdt 1  
Florence, SC, KFLO, RNAV (GPS) RWY 9,  
Amdt 1  
Florence, SC, KFLO, RNAV (GPS) RWY 19,  
Amdt 1  
Florence, SC, KFLO, RNAV (GPS) RWY 27,  
Amdt 1  
Sumter, SC, KSMS, ILS OR LOC RWY 23,  
Amdt 2  
Sumter, SC, KSMS, Takeoff Minimums and  
Obstacle DP, Amdt 1A  
Seattle, WA, KBFI, RNAV (GPS) Y RWY 14R,  
Orig  
Seattle, WA, KBFI, RNAV (GPS) Y RWY 14R,  
Amdt 1A, CANCELED  
Seattle, WA, KBFI, RNAV (GPS) Y RWY 32L,  
Orig  
Seattle, WA, KBFI, RNAV (RNP) Z RWY 14R,  
Orig  
Seattle, WA, KBFI, RNAV (RNP) Z RWY 14R,  
Amdt 1A, CANCELED  
Seattle, WA, KBFI, RNAV (RNP) Z RWY 32L,  
Orig  
*Rescinded:* On June 21, 2023 (88 FR  
40081), the FAA published an Amendment  
in Docket No. 31490, Amdt No. 4063, to part  
97 of the Federal Aviation Regulations under  
§§ 97.23, 97.29, and 97.33. The following  
entries for Northway, AK, San Francisco, CA,  
and Cross Keys, NJ, effective August 10,  
2023, are hereby rescinded in their entirety:  
Northway, AK, PAOR, RNAV (GPS) RWY 6,  
Amdt 1  
Northway, AK, PAOR, RNAV (GPS) RWY 24,  
Amdt 2  
San Francisco, CA, KSFO, GLS RWY 19L,  
Amdt 1  
San Francisco, CA, KSFO, GLS RWY 19R,  
Amdt 1  
Cross Keys, NJ, 17N, VOR OR GPS RWY 9,  
Amdt 6B, CANCELED

[FR Doc. 2023–14932 Filed 7–13–23; 8:45 am]

**BILLING CODE 4910–13–P**

## FEDERAL TRADE COMMISSION

### 16 CFR Parts 0, 1, 2, 3 and 4

#### Rules of Practice

In rule document 2023–12630 beginning on page 42872 in the issue of Wednesday, July 5, 2023, make the following corrections:

On page 42872, in the third column, under **DATES**, in the first and fourth lines “June 5, 2023” should read “July 5, 2023”.

[FR Doc. C1–2023–12630 Filed 7–13–23; 8:45 am]

**BILLING CODE 0099–10–D**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### 21 CFR Chapter I

[Docket No. FDA–2023–N–0963]

#### Nomenclature Change for Dockets Management; Technical Amendment

**AGENCY:** Food and Drug Administration, Department of Health and Human Services (HHS).

**ACTION:** Final rule; technical amendment.

**SUMMARY:** The Food and Drug Administration (FDA) is amending its regulations to reflect a change in the name of Division of Dockets Management to Dockets Management Staff and information regarding copies. This action is editorial in nature and is intended to improve the accuracy of the Agency’s regulations.

**DATES:** This rule is effective July 14, 2023.

**FOR FURTHER INFORMATION CONTACT:** Karen Malvin, Dockets Management Staff, Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

**SUPPLEMENTARY INFORMATION:** FDA is amending 21 CFR chapter I to update Dockets Management Staff’s name change and information regarding copies.

Publication of this document constitutes final action on the changes under the Administrative Procedure Act (5 U.S.C. 553). FDA has determined that notice and public comment are unnecessary because this amendment to the regulations provides only a technical change to update the organizational information for Dockets Management Staff.

#### List of Subjects

##### 21 CFR Part 3

Administrative practice and procedure, Biologics, Drugs, Medical devices.

##### 21 CFR Part 5

Authority delegations (Government agencies), Imports, Organization and functions (Government agencies).

##### 21 CFR Part 7

Administrative practice and procedure, Consumer protection, Reporting and recordkeeping requirements.

##### 21 CFR Part 10

Administrative practice and procedure, News media.

##### 21 CFR Parts 12, 13, and 15

Administrative practice and procedure.

##### 21 CFR Part 14

Administrative practice and procedure, Advisory committees, Color additives, Drugs, Radiation protection.

##### 21 CFR Part 17

Administrative practice and procedure, Penalties.

##### 21 CFR Part 20

Confidential business information, Courts, Freedom of information, Government employees.

##### 21 CFR Part 25

Environmental impact statements, Foreign relations, Reporting and recordkeeping requirements.

##### 21 CFR Part 60

Administrative practice and procedure, Drugs, Food additives, Inventions and patents, Medical devices, Reporting and recordkeeping requirements.

##### 21 CFR Part 100

Administrative practice and procedure, Food labeling, Food packaging, Foods, Intergovernmental relations.

##### 21 CFR Part 101

Food labeling, Nutrition, Reporting and recordkeeping requirements.

##### 21 CFR Part 109

Food packaging, Foods, Polychlorinated biphenyls (PCBs).

##### 21 CFR Part 165

Beverages, Bottled water, Food grades and standards.

##### 21 CFR Part 170

Administrative practice and procedure, Food additives, Reporting and recordkeeping requirements.

##### 21 CFR Part 184

Food additives.

##### 21 CFR Part 201

Drugs, Labeling, Reporting and recordkeeping requirements.

##### 21 CFR Part 312

Drugs, Exports, Imports, Investigations, Labeling, Medical research, Reporting and recordkeeping requirements, Safety.

##### 21 CFR Part 314

Administrative practice and procedure, Confidential business